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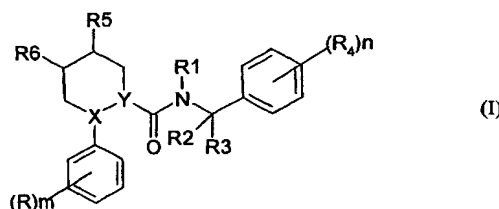
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(54) Title: PIPERIDYLCARBOXAMIDE DERIVATIVES AND THEIR USE IN THE TREATMENT OF TACHYKINIM-MEDIATED DISEASES



(57) Abstract: The present invention relates to piperidine derivatives of formula (I) wherein R represents halogen or C<sub>1-4</sub>alkyl; R<sub>1</sub> represents hydrogen or C<sub>1-4</sub> alkyl; R<sub>2</sub> represents hydrogen, C<sub>1-4</sub> alkyl or R<sub>2</sub> together with R<sub>3</sub> represents C<sub>3-7</sub> cycloalkyl; R<sub>3</sub> represents hydrogen, C<sub>1-4</sub> alkyl, C<sub>3-7</sub> cycloalkyl or C<sub>3-6</sub> alkenyl; or R<sub>1</sub> and R<sub>3</sub> together with nitrogen and carbon atom to which they are attached respectively represent a 5 to 6 membered heterocyclic group; R<sub>4</sub> represents trifluoromethyl, C<sub>1-4</sub> alkyl, C<sub>1-4</sub> alkoxy, trifluoromethoxy or halogen; R is hydrogen and R is NR<sub>7</sub>R<sub>8</sub> or R<sub>5</sub> is NR<sub>8</sub>R<sub>9</sub> and R<sub>6</sub> is hydrogen; R<sub>7</sub> represents hydrogen or C<sub>1-4</sub> alkyl or R<sub>7</sub> and R<sub>8</sub> together with nitrogen to which they are attached are a saturated 5 to 7 membered heterocyclic group containing oxygen; R<sub>8</sub> represents hydrogen, phenyl, C<sub>3-7</sub> cycloalkyl, (CH<sub>2</sub>)<sub>p</sub>C(O)NR<sub>10</sub>R<sub>11</sub>, a saturated 5 to 7 membered heterocyclic group containing 1 to 3 heteroatoms selected from oxygen, sulphur and nitrogen and optionally substituted by C<sub>1-4</sub> alkyl, S(O)<sub>2</sub>C<sub>1-4</sub> alkyl or C(O) C<sub>1-4</sub> alkyl, a 5 membered heteroaryl group containing 1 to 3 heteroatoms selected from oxygen, sulphur and nitrogen and optionally substituted by C<sub>1-4</sub> alkyl S(O)<sub>2</sub>C<sub>1-4</sub>alkyl or C(O) C<sub>1-4</sub> alkyl or R<sub>8</sub> represents a 6 membered heteroaryl group containing 1 to 3 nitrogen atoms and optionally substituted by C<sub>1-4</sub>alkyl, S(O)<sub>2</sub>C<sub>1-4</sub> alkyl or C(O) C<sub>1-4</sub>alkyl; or R<sub>8</sub> is a C<sub>1-6</sub> alkyl group optionally substituted by one or two groups selected from fluorine, phenyl(optionally substituted by C<sub>1-4</sub> alkyl, C(O) C<sub>1-4</sub> alkyl or halogen), =O, C<sub>3-7</sub>cycloalkyl, hydroxy, amino, dimethylamino, aminocarbonyl, C<sub>1-4</sub>alkoxy or trifluoromethyl; R<sub>9</sub> is hydrogen, C<sub>1-4</sub> alkyl or R<sub>9</sub> and R<sub>8</sub> together with nitrogen to which they are attached are a 5 to 7 membered heterocyclic group optionally containing another heteroatom selected from oxygen, sulphur and nitrogen and optionally substituted by one or two groups selected from C<sub>1-4</sub> alkyl, =O, S(O)<sub>2</sub>C<sub>1-4</sub> alkyl, C(O) C<sub>3-7</sub>cycloalkyl or C(O) C<sub>1-4</sub> alkyl; R<sub>10</sub> and R<sub>11</sub> are independently hydrogen or C<sub>1-4</sub> alkyl group; X represents a nitrogen atom and Y is CH or X represents CH and Y is nitrogen; m is zero or an integer from 1 to 3; n is an integer from 1 to 3; p is zero, 1 or 2; and pharmaceutically acceptable salts and solvates thereof; the process for their preparation and their use in the treatment of conditions mediated by tachykinins.